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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/568,226	02/14/2006	Yasuo Kunugiza	GRT/423-72	6159	
23117 NIXON & VA	7590 07/21/200 NDERHYE, PC	8	EXAM	EXAMINER	
901 NORTH GLEBE ROAD, 11TH FLOOR			EPPS FORD, JANET L		
ARLINGTON	, VA 22203		ART UNIT	PAPER NUMBER	
			1633	•	
			MAIL DATE	DELIVERY MODE	
			07/21/2008	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Application No. Applicant(s) 10/568,226 KUNUGIZA ET AL.

Office Action Summary	Examiner	Art Unit					
	Janet L. Epps-Ford	1633					
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply							
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA Extensions of time may be available under the provisions of 37 CFR 1.13 after SSI (6) MONTHS from the mailing date of this communication. If NO period for reply is specified above, the maximum statutory period we Failure to reply within the set or extended period for reply with y statute, Any reply received by the Office later than three months after the mailing carried patient term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tin viil apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	N. nely filed the mailing date of this o D (35 U.S.C. § 133).	,				
Status							
1) Responsive to communication(s) filed on 17 Ap	oril 2008.						
2a) This action is FINAL. 2b) ☐ This	action is non-final.						
 Since this application is in condition for allowar 	nce except for formal matters, pro	secution as to the	merits is				
closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.							
Disposition of Claims							
4) ☐ Claim(s) 1-18 and 20 is/are pending in the appi 4a) Of the above claim(s) is/are withdraw 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 1-9.11-18 and 20 is/are rejected. 7) ☐ Claim(s) 10 is/are objected to. 8) ☐ Claim(s) are subject to restriction and/or	vn from consideration.						
Application Papers							
9) The specification is objected to by the Examine: 10) The drawing(s) filed on is/are: a) acc Applicant may not request that any objection to the c Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	epted or b) objected to by the I drawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). jected to. See 37 Cl					
Priority under 35 U.S.C. § 119							
12) 🖾 Acknowledgment is made of a claim for foreign a) 🖾 All b) 🗀 Some * c) 🗀 None of: 1. 🖾 Certified copies of the priority documents 2. ☐ Certified copies of the priority documents 3. ☐ Copies of the certified copies of the prior application from the International Bureau * See the attached detailed Office action for a list of	s have been received. s have been received in Applicati ity documents have been receive I (PCT Rule 17.2(a)).	on No ed in this National	Stage				
Attachment(s)							
1) Notice of References Cited (PTO-892) 2) Notice of Professorson's Patent Drawing Review (PTO-948)	4) Interview Summary Paper No(s)/Mail Da						

3) Information Disclosure Statement(s) (PTO/S6/08) Paper No(s)/Mail Date 2-16-06.

5) Notice of Informal Patent Application 6) Other: _____.

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DETAILED ACTION

Election/Restrictions

1. Applicant's election with traverse of Group 1 (claims 1-18), drawn to a staple oligonucleotide of formula SEQ ID NO: 1, in the reply filed on 4-17-2008 is acknowledged. The traversal is on the ground(s) that the examination of Groups I to III would not constitute a serious burden. This is not found persuasive because although SEQ ID NOS: 1-3 share some structural similarities, nevertheless overall each claimed staple oligonucleotide comprise a different nucleotide sequence and therefore require a separate search and consideration of the prior art.

The requirement is still deemed proper and is therefore made FINAL.

 Groups II-III, drawn to SEQ ID NO: 2-3 are withdrawn from further consideration pursuant to 37 CFR 1.142(b), as being drawn to a nonelected invention, there being no allowable generic or linking claim. Applicant timely traversed the restriction (election) requirement in the reply filed on 4-17-2008.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- Claims 1-, 6, 8-9, and 11-18 are rejected under 35 U.S.C. 102(b) as being anticipated by Vasseur et al. (WO 94/23026).
- Claim 1 recites the following:

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1. (original) A staple oligonucleotide which is a single-stranded oligonucleotide comprising a 5'-end sequence, an intermediate sequence and a 3'-end sequence, the 5'-end sequence having a reverse complementarity with the intermediate sequence, the 3'-end sequence having a reverse complementarity with the intermediate sequence and the intermediate sequence having loops at both ends, the loops each comprising three to ten nucleotides and not forming a complementary bond intermolecularly.

- 6. (previously presented) The staple oligonucleotide according to claim 1, wherein the loops each comprise 4 to 6 nucleotides in length.
- 8. (previously presented) The staple oligonucleotide according to claim 1, wherein the oligonucleotide is a DNA or a DNA derivative.
- 9. (previously presented) The staple oligonucleotide according to claim 1, whose phosphate groups are not phosphorothioated.

The following structure is disclosed in Vasseur et al. in Figures 2A and 4a. This structure meets all the limitations recited in instant claims 1, 6, and 8-9, particularly wherein the staple oligonucleotide comprises a 3'-end and a 5'end comprising reverse complementarity, and 4-nucleotide loops on both ends. The disclosed oligonucleotide is a DNA derivative, and there is no indication that the compounds are phosphorothioated, see Figure 1.

In regards to the intended use limitations recited in instant claims 11-18, particularly wherein the generically claimed staple oligonucleotide is disclosed as a "medicament," absent evidence to the contrary, since the prior art describes the general structure recited in instant claim 1, the prior art structure would have to also meet the intended use limitations recited in the instant claims, and thus anticipate the claimed invention.

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 Claims 1-9, 11-18 are rejected under 35 U.S.C. 102 (b) as being anticipated by Blumenfeld et al. (WO 9219731 A1).

Figure 5B of this reference discloses the following:

FIGURE 5-B.

This compound comprises 48 nucleotides in length, and thus falls within the range of 42 to 54 nucleotides as recited in the instant claims, and further comprises loops of 4 to 6 nucleotides in length.

In regards to the intended use limitations recited in instant claims 11-18, particularly wherein the generically claimed staple oligonucleotide is disclosed as a "medicament," absent evidence to the contrary, since the prior art describes the general structure recited in instant claim 1, the prior art structure would have to also meet the intended use limitations recited in the instant claims, and thus anticipate the claimed invention.

- Claims 1-9, 11-18 and 20 are rejected under 35 U.S.C. 102(b) as being anticipated by Ahn et al.
- Ahn et al. describe decoy oligonucleotides which specifically bind the transcription factor E2F, see Figure 1, CDODN, MODN is the mutated control.

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A A GUARIAGE GEDAARC GGATCOCTTTE GEGETATTOE A A
                    A _CUTTATEGEGETETOCCTAGOCAAAGGGGGATAACG _ A
                    * GCAATAAAICGAAACGCATCCGHHEGAIHTATREE * A
A GGTTATHIAGCIHTGCCTAGGCAAAGCTAAATAACG , A
MODN
Figure 7. Structures and sequences of the decay ODNs used in this study. The two E25 recognition sequences are undertised so cach mechanide. CD-E25 ODN continue two binding sites for E25 in its clear region.
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10. Ahn et al. demonstrated that the E2F decoy, CD-ODN inhibited the growth of vascular smooth muscle cells in vitro, and further demonstrated a significant reduction of neointimal formation in a dose dependent manner comprising the administration of CD-ODN in vivo.

Allowable Subject Matter

11. Claim 10 is objected to as being dependent upon a rejected base claim, but would be allowable if rewritten to recite the elected subject matter, and further rewritten in independent form including all of the limitations of the base claim and any intervening claims.

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12. Any inquiry concerning this communication or earlier communications from the

examiner should be directed to Janet L. Epps-Ford whose telephone number is 571-

272-0757. The examiner can normally be reached on M-F, 10:00 AM through 6:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Joseph Woitach can be reached on 571-272-0739. The fax phone number

for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the

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system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Janet L. Epps-Ford/

Primary Examiner, Art Unit 1633

/J. L. E./

Primary Examiner, Art Unit 1633